Application No.: 09/509648 Docket No.: JJJ-P01-569

# **AMENDMENTS TO THE CLAIMS**

 (Currently amended) A method for potentiating a morphogen activity in a neuron, comprising administering to a mammal contacting the neuron with a composition, the composition comprising a molecule which: that

- (a) is a neuropoietic cytokine antagonist, a retinoid antagonist, or a cAMPdependent messenger pathway inhibitor; and
- (b) overcomes morphogen-inhibition of the morphogen activity in vitro; thereby potentiating the morphogen activity in a neuron.
- (Currently amended) A method for promoting neuronal cell growth, comprising administering to a mammal contacting a neuron with a composition, the composition comprising a molecule which: that
  - (a) is a neuropoietic cytokine antagonist, a retinoid antagonist, or cAMPdependent messenger pathway inhibitor; and
  - (b) overcomes morphogen inhibition, of so as to potentiate-growth-promoting effects of endogenous morphogens in vitro;

thereby promoting neuronal cell growth.

### 3 - 4. (Canceled)

- 5. (Previously presented) The method of claim 1, wherein said morphogen activity is endogenous.
- 6. (**Previously presented**) The method of claim 1, wherein said morphogen activity is the result of an exogenously provided morphogen.
- 7. (Currently amended) The method of claim 42, wherein said composition further comprises a morphogen.
- 8. (Currently amended) The method of claim 3-or 41 or 2, wherein said disorder neuron is injured by Alzheimer's disease, Parkinson's disease, Huntington's disease, senile dementia, alcohol-induced dementia, or stroke.

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## 9-10. (Canceled)

11. (Currently amended) The method of claim 10-1 or 2, wherein said neuropoetic neuropoietic cytokine antagonist is an LIF (Leukemia-Inhibitory Factor) antagonist or a CNTF (Ciliary Neurotrophic Factor) antagonist.

12. (Currently amended) The method of claim 11, wherein said neuropoetic cytokine antagonist is a LIF (Leukemia-Inhibitory Factor) antagonist which is a monoclonal antibody to a gp130 protein.

## 13-15. (Canceled)

- 16. (Previously presented) The method of claim 7, wherein said morphogen comprises an amino acid sequence selected from a sequence:
  - (a) having at least 70% homology with the C-terminal seven-cysteine skeleton of human OP-1 (Osteogenic Protein 1), residues 330-431 of SEQ ID NO: 2;
  - (b) having greater than 60% amino acid sequence identity with said C-terminal seven-cysteine skeleton of human OP-1;
  - (c) defined by Generic Sequence 7, SEQ ID NO: 4;
  - (d) defined by Generic Sequence 8, SEQ ID NO: 5;
  - (e) defined by Generic Sequence 9, SEQ ID NO: 6;
  - (f) defined by Generic Sequence 10, SEQ ID NO: 7; or
  - (g) defined by OPX, SEQ ID NO: 3.
- 17. (Previously presented) The method of claim 7, wherein said morphogen is human OP-1 (Osteogenic Protein 1), mouse OP-1, human OP-2 (Osteogenic Protein 2), mouse OP-2, 60A, GDF-1 (Growth/Differentiation Factor-1), BMP2A (Bone Morphogenesis Protein 2A), BMP2B (Bone Morphogenesis Protein 2B), DPP (Decapentaplegic), Vgl, Vgr-1 (Vg1-related sequence), BMP3 (Bone Morphogenesis Protein 3), BMP5 (Bone Morphogenesis Protein 5), or BMP6 (Bone Morphogenesis Protein 6).
- 18. (Previously presented) The method of claim 7, wherein said morphogen is OP-1.

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19. (Currently amended) The method of claim 1, wherein the molecule neuropoietic cytokine antagonist binds an endogenous ligand for a cytokine receptor-or a retinoid receptor.

# 20 - 21. (Canceled)

22. (Currently amended) The method of claim 191 or 2, wherein the molecule which binds an endogenous ligand for a retinoid receptor retinoid antagonist is a retinoic acid receptor or retinoid X receptor.

## 23 - 24. (Canceled)

- 25. (Currently amended) The method of claim 241, wherein said cAMP-dependent messenger pathway inhibitor comprises a protein kinase A inhibitor.
- 26. (**Previously presented**) The method of claim 25, wherein said protein kinase A inhibitor is (2-p-bromocynnamylaminoethyl)-5-isoquinolinesulfonamide, an enantiomer of dibutyryl cAMP, or an enantiomer of cAMP.

## 27 - 32. (Canceled)

- 33. (New) The method of claim 1, wherein the retinoid antagonist binds an endogenous ligand for a retinoid receptor.
- 34. (New) The method of claim 1, wherein said morphogen activity is activity to stimulate dendritic growth.
- 35. (New) The method of claim 1, wherein said morphogen activity is activity of OP-1.
- 36. (New) The method of claim 2, wherein said neuronal cell growth is dendritic growth.
- 37. (New) The method of claim 1, 2, 34 or 36, wherein said neuron is a sympathetic neuron.
- 38. (New) The method of claim 11, wherein said CNTF inhibitor is phosphatidylinositol-specific phospholipase C.